GUAIFENESIN- guaifenesin solution
Cardinal Health 107, LLC

GUAIFENESIN

Expectorant
SUGAR FREE / ALCOHOL FREE

DESCRIPTION

Each 5 mL (1 teaspoonful) contains:
Guaifenesin 100 mg

Inactive Ingredients

Acesulfame K, citric acid, FD&C Green No. 3, FD&C Red No. 40, flavoring, hydroxyethylcellulose, purified water, sodium benzoate and sodium citrate.

Sodium Content: 4 mg/5 mL

USES

Helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive.

WARNINGS

Ask a doctor before use if you have

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.
- you are hypersensitive to any of the ingredients.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Professional Note

Guaifenesin has been shown to produce a color interference with certain clinical laboratory determinations of 5-hydroxyindoleacetic acid (5-HIAA) and vanillylmandelic

acid (VMA).

DIRECTIONS

Follow dosage below or use as directed by a physician.

- do not take more than 6 doses in any 24-hour period.

age	dose
adults and children 12 years and over	10 to 20 mL (2 to 4 teaspoonfuls) every 4 hours
children 6 years to under 12 years	5 to 10 mL (1 to 2 teaspoonfuls) every 4 hours
children 2 to under 6 years of age	2.5 to 5 mL (½ to 1 teaspoonful) every 4 hours
children under 2 years of age	consult a physician

Guaifenesin Oral Solution USP is a red, raspberry flavored solution supplied in the following oral dosage forms:

Overbagged with 5 x 5 mL unit dose cups in each bag, NDC 55154-9450-5

Overbagged with 5 x 10 mL unit dose cups in each bag, NDC 55154-5780-5

WARNING: These Unit Dose packages are not child resistant and are Intended for Institutional Use Only. Keep this and all drugs out of the reach of children.

Distributed By:

Cardinal Health

Dublin, OH 43017

L54670551025

L53963871025

R04/05 F1744050221 F1744100221 F1744150221

STORAGE

Keep tightly closed. Store at controlled room temperature, 20°-25°C (68°-77°F). [See USP] Protect from light.

Principal Display Panel

NDC 55154-9450-5

GUAIFENESIN ORAL SOLUTION USP

100 mg/5mL

5 CUPS

N108

NDC 55154-9450-5

GUAIFENESIN ORAL SOLUTION USP

100 mg/5 mL

5 CUPS

EXPECTORANT

Sugar Free / Alcohol Free
Delivers 5 mL

See product insert for prescribing information, precautions and warnings.

STORAGE: Store at 20° to 25° C (68° to 77° F)
[See USP Controlled Room Temperature]

WARNING: This Unit Dose package is not child resistant and is Intended for Institutional Use Only.
Keep this and all drugs out of the reach of children.

PAI Pharma
Greenville, SC 29605
1-800-845-8210
www.paipharma.com
DURA-DOSE®

Distributed by Cardinal Health
Dublin, OH 43017

L54670551025

Lot: Exp:

Principal Display Panel

NDC 55154-5780-5

GUAIFENESIN ORAL SOLUTION USP

200 mg/10 mL

5 CUPS

I102

NDC 55154-5780-5

GUAIFENESIN ORAL SOLUTION USP
200 mg/10 mL

5 CUPS

EXPECTORANT

Sugar Free / Alcohol Free

Delivers 10 mL

See product insert for prescribing information, precautions and warnings.

STORAGE: Store at 20° to 25° C (68° to 77° F)
[See USP Controlled Room Temperature]

WARNING: This Unit Dose package is not child resistant and is Intended for Institutional Use Only.
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GREENVILLE, SC 29605
1-800 845-8210
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DURA-DOSE®

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Lot: Exp:

GUAIFENESIN
guaifenesin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55154-9450(NDC:0121-1744)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136Y38GY5)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	red	**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-9450-5	5 in 1 BAG	09/01/2002	
1		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/01/2002	

GUAIFENESIN

guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55154-5780(NDC:0121-1488)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136Y38GY5)	
WATER (UNII: 059QF0KO0R)	
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SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	red	**Score**	
Shape		**Size**	
Flavor	RASPBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-5780-5	5 in 1 BAG	09/01/2002	
1		10 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		

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OTC Monograph Drug	M012	09/01/2002	

Labeler - Cardinal Health 107, LLC (118546603)

Revised: 6/2026 Cardinal Health 107, LLC